

TRANSPORTATION OF BIOLOGICAL MATTER TO A TARGET SITE IN A CLOSED VOLUME FOR TRANSPLANTATION PURPOSES

FIELD OF THE INVENTION

The invention is in the field of transporting biological matter to a target site
5 in a closed volume for transplantation purposes.

BACKGROUND OF THE INVENTION

In commonly assigned WO99/18872 entitled *"Method for Depositing a
Flattened Droplet on a Surface and Apparatus Therefor, and a Pump Therefor"*, the
contents of which are incorporated herein by reference, there is illustrated and
10 described a flattened droplet type IVF-ET procedure in which preferably a single
embryo is accurately deposited on an embryo recipient subject's endometrium. One
advantage of the flattened droplet approach is that it employs only about 0.3-2 μ l
culture medium to transport an embryo to a target site as opposed to 20-40 μ l
culture medium in the conventional IVF-ET injection approach but surprisingly,
15 even this microvolume may increase a typical 3-4 Inches of Water uterine
prevailing pressure by 1-3 Inches of Water. Another advantage of the flattened
droplet approach is that an embryo is subjected to a far smaller pressure than in
comparison to the conventional IVF-ET injection approach. Clinical trials have
demonstrated that the flattened droplet approach leads to a considerably higher
20 percentage of successful pregnancies in comparison to the conventional IVF-ET
injection approach. However, this notwithstanding, the flattened droplet IVF-ET
approach still suffers from technical faults, for example, kinking of a delivery
catheter, blockage of its distal end, and the like, which in all likelihood lead to an
unsuccessful IVF-ET procedure.

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SUMMARY OF THE INVENTION

The present invention is directed toward the transportation of biological matter to a target site in a closed volume for transplantation purposes. The present invention involves provisioning the aforementioned WO99/18872's flattened
5 droplet delivery apparatus with a pressure sensor for real time monitoring of the prevailing pressure in its transfer tube. The prevailing pressure can be displayed on a monitor particularly during the actual transfer of an embryo bearing culture medium microvolume to an embryo recipient subject's endometrium such that an occurrence of one or more of the aforelisted technical faults would be immediately
10 apparent to a trained practitioner who may then take appropriate corrective action. Pattern recognition functionality can be readily applied to a pressure waveform of the prevailing pressure within a transfer tube for detecting an occurrence of one or more of the aforelisted technical faults by virtue of each technical fault having a uniquely identifiable fault signature, thereby enabling the automatic issuance of a
15 visual and/or aural alarm in the case of such an occurrence. To largely negate the aforesaid typical 1-3 Inches of Water prevailing uterine pressure increase during a flattened droplet type IVF-ET procedure, the delivery catheter for delivering the embryo bearing culture medium microvolume to an embryo recipient subject's uterus is preferably introduced through an extruded guide catheter designed to
20 slidably support the threading of the delivery catheter therethrough, and enable concurrent fluid drainage from her uterus to lower her prevailing uterine pressure.

The present invention also enables the transportation of a series of biological matter bearing flattened droplets for effecting cell based therapy procedures rather than the presently suggested injection approach as described in an
25 on-line article entitled "Stem Cells: A primer" as retrieved from the National Institute of Health (NIH)'s website on April 12, 2003 <URL: www.nih.gov/news/stemcell/primer.htm>. Cell based therapies are now being hypothesized to treat a wide range of diseases including *inter alia* Parkinson's disease, diabetes, traumatic spinal cord injury, heart disease, vision and hearing
30 loss, and the like. In this context, biological matter may be in the form of complete

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cells, for example, stem cells, germ cells, and the like, and also cell components, for example, DNA, RNA, and the like. It is believed that the flattened droplet approach in comparison to the conventional injection approach will be particularly advantageous for cell based therapy procedures by virtue of the fact that the aforesaid advantages of the former approach will be even more beneficial due to cell based therapy procedures requiring the transplantation of far greater quantities of biological matter than preferably a single embryo for an IVF-ET procedure, and the target sites of cell based therapy procedures are in closed volumes far smaller than a human uterus and therefore more susceptible to increases in prevailing pressures.

BRIEF DESCRIPTION OF DRAWINGS

In order to understand the invention and to see how it may be carried out in practice, preferred embodiments will now be described, by way of non-limiting examples only, with reference to the accompanying drawings, in which:

Fig. 1 is a pictorial representation of apparatus for effecting a flattened droplet type IVF-ET procedure in accordance with the present invention;

Fig. 2 is a transverse cross section view of a guide catheter with a delivery catheter threaded therethrough for use in a flattened droplet type IVF-ET procedure;

Figs. 3A-3F are pictorial representations showing the transport of an embryo bearing culture medium microvolume to an embryo recipient subject's endometrium during a flattened droplet type IVF-ET procedure;

Fig. 4 is a graph showing the pressure waveform of the prevailing pressure in the transfer tube of the apparatus of Figure 1 during a flattened droplet type IVF-ET procedure together with fault signatures of three potential technical faults which may occur during such a procedure;

Fig. 5 is a pictorial representation of apparatus for transporting biological matter for a cell based therapy procedure in accordance with the present invention;

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Figs. 6A to 6C are pictorial representations showing the operation of the apparatus of Figure 5;

Fig. 7 is a graph showing the prevailing pressure within the transfer tube of the apparatus of Figure 5 during a cell based therapy procedure; and

5 Fig. 8 is a pictorial representation showing the fusing together of a number of stem cell bearing flattened droplets into a large drop.

DESCRIPTION OF PREFERRED EMBODIMENTS

Figure 1 shows apparatus 1 for effecting an improved flattened droplet type IVF-ET procedure. Apparatus 1 includes a suction control unit 2 typically
10 permanently located in a laboratory for the preparation of an embryo bearing delivery catheter (constituting a narrow bore transfer tube) 3, a transfer control unit 4 typically permanently located in a treatment room where IVF-ET procedures are carried out and a portable casing 6 for consecutive connection to the suction control unit 2 and the transfer control unit 4 by means of connectors 7 and 8. Insertion of
15 the delivery catheter 3 into an embryo recipient subject's uterus is preferably through an extruded guide catheter 9 having, say, four to eight, longitudinally directed supports 11 whose longitudinally directed curved inner facing surfaces 12 define an imaginary circular in the transverse cross section view of Figure 2 with a diameter slightly larger than the delivery catheter's diameter D whereby the delivery
20 catheter 3 can be readily slidingly threaded therethrough. The use of the guide catheter 9 enables fluid drainage from an embryo recipient subject's uterus to avoid a significant increase in her uterine pressure which may militate against a successful IVF-ET procedure.

The casing 6 includes a pneumatic system 13 which is permanently
25 connected to the delivery catheter 3 during an entire IVF-ET procedure via suitable air tubing 14, and a receptacle 16 for accommodating the delivery catheter 3 during the transport of the casing 6 from a laboratory to a treatment room. The pneumatic system 13 includes a microvolume pump (not shown), for example, as described with reference to WO99/18872's Figures 3-5, and a pressure sensor 17 for

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monitoring the prevailing pressure within the delivery catheter 3 for display as a Single Flattened Droplet Pressure Waveform (SFDPW) on a computer 18. The pressure sensor 17 is operative over 0-1 psi a pressure range and has an about $\pm 1\%$ pressure sensitivity. IC Sensor's Model 1210 pressure sensor would be suitable for use as pressure sensor 17. The computer 18 can be programmed with pattern recognition functionality 19 to automatically issue visual and/or aural alarms on detection of an occurrence of one or more fault signatures in the SFDPW as described hereinbelow with reference to Figure 4.

The pneumatic system 13 is under the control of a control mechanism 21 including a control pad 22 for controlling the suction control unit 2 for initiating a user controlled suction mode to load the delivery catheter 3 with an embryo bearing culture medium microvolume and a foot pedal 23 for controlling the transfer control unit 4 for initiating a user initiated automated delivery mode for depositing an embryo bearing flattened droplet on an embryo recipient subject's endometrium. The control pad 22 has an upstroke control 22A for drawing an incoming flow of displacement gas into the pneumatic system 13 from the delivery catheter 3, a downstroke control 22B for issuing an outgoing flow of displacement gas from the pneumatic system 13 into the delivery catheter 3, and optionally a speed control 22C for controlling the flow rate of the displacement gas either from or into the delivery catheter 3. The suction control unit 2 is also provided with a reset button 26 for priming the pneumatic system 13 for a pre-suction mode of issuing an outgoing flow of displacement gas as indicated by a READY indicator light 27 prior to the loading of the delivery catheter 3. The different stages of the automatic delivery mode are indicated by a READY indicator light 28, a GO indicator light 29 and a DONE indicator light 31.

For the sake of conciseness, the loading of the delivery catheter 3 with one or more embryo(s) in accordance with the procedure described with reference to WO99/18872's Figures 2A-2E is not repeated here. The depositing of an embryo bearing flattened droplet on an embryo recipient subject's endometrium S is now

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described with reference to the steps shown in Figures 3A-3F and SFDPW shown in Figure 4.

The delivery catheter's distal end 3A is laid on an embryo recipient subject's endometrium S (see Fig. 3A) whereupon a first depression on the foot pedal 23 causes the READY indicator light 28 to be lit indicating that the automatic delivery mode can be initiated. Thereafter, a second depression on the foot pedal 23 causes the GO indicator light 29 to be lit indicating that an outgoing flow of displacement gas is displacing the embryo bearing culture medium microvolume towards the distal end 3A (see Fig. 3B). The outgoing flow of displacement gas causes a concave shaped meniscus to be slowly formed which increases in size until it suddenly ruptures whereby most of the embryo bearing culture medium microvolume is discharged as a droplet D on the surface S (see Figs. 3C and 3D). The discharge is accompanied by blowing miniscule air bubbles B into the droplet D for frothing it and thereby considerably widening its projected surface area on the embryo recipient subject's endometrium S to form the flattened droplet F whose shape is maintained by its prevailing surface tension with the embryo recipient subject's endometrium S (see Fig. 3E).

The GO indicator light 29 is then extinguished indicating that the practitioner should slightly withdraw the delivery catheter 3 so as to detach its distal end 3A from the droplet F whilst at the same time there is a slow discharge of displacement gas. Withdrawal is limited to between about 10-15 mm such that the distal end 3A still lies along the embryo recipient subject's endometrium S. And finally, a further outgoing pulse of displacement gas is provided to clean the distal end 3A of any remaining culture medium (see Fig. 3F). The DONE indicator light 31 is then lit to indicate that the delivery catheter 3 can be completely removed from the embryo recipient subject's uterus.

Figure 4 also shows three fault signatures FS1, FS2 and FS3 superimposed on the SFDPW of technical faults which may occur during a flattened droplet type IVF-ET procedure as follows:

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A fault signature FS1 appears as a pressure drop at the beginning of the SFDPW during an initial issue of outgoing flow of displacement gas to outwardly displace the embryo bearing culture medium microvolume along the delivery catheter 3. A fault signature FS1 is indicative that the pneumatic system 13 is not hermetically sealed and that suitable action be taken to hermetically seal the pneumatic system 13 before a flattened droplet type IVF-ET procedure can be continued.

A fault signature FS2 appears as a pressure increase beyond a predetermined maximum pressure, say, 12 Inches of Water for an IVF-ET procedure. A fault signature FS2 is indicative of either a kink along the delivery catheter 3 or a blockage at its distal end 3A due to soft tissue or blood. A fault signature FS2 requires that a practitioner attempt to unkink the delivery catheter 3 or unblock its distal end 3A by gently manipulate the delivery catheter 3 to and fro or up and down.

A fault signature FS3 appears as a pressure increase towards the end of the SFDPW after the issue of the further outgoing pulse of displacement gas denoted 3F in Figure 4. The pressure increase appears instead of the prevailing pressure in the delivery catheter 3 dropping down to a typical uterine prevailing pressure of about 3-4 Inches of Water. A fault signature FS3 is indicative of the back flow of a flattened droplet F possibly including the embryo E into the delivery catheter 3. A fault signature FS3 requires that a practitioner issue a second outgoing pulse of displacement gas.

Figure 5 shows apparatus 51 for automatically transporting a series of biological matter bearing flattened droplets to a target site for, say, a stem cell therapy procedure. The apparatus 51 includes a pneumatic system 52 under the control of a computer (constituting a control mechanism) 53 including the pattern recognition functionality 19 programmed for detecting the fault signatures FS1, FS2 and FS3, and issuing suitable alarms, and a tubing set 54 designed for single or multiple use. The pneumatic system 52 includes a microvolume pump 56, for example, as described with reference to WO99/18872's Figures 3-5.

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The tubing set 54 includes a transfer tube 57 having a proximal end 57A, a distal end 57B, and a pair of tube segments 57C and 57D; a two-way valve 58 under the control of the computer 53 and having three ports 58A, 58B and 58C; an uptake catheter 59 having a proximal end 59A and a distal end 59B; and a delivery catheter 61 having a proximal end 61A and a distal end 61B. The prevailing pressure in the transfer tube 57 is monitored by a pressure sensor 62 hermetically connected to the tube segment 57C and connected to the computer 53 for real time feedback purposes and for displaying a Multiple Flattened Droplet Pressure Waveform (MFDPW) arising from a stem cell therapy procedure. The transfer tube 57 can be selectively vented to atmospheric pressure P_0 by a normally closed (NC) venting valve 63 hermetically connected to the tube segment 57D and under the control of the computer 53.

The computer 53 also receives closed loop feedback from an electro-optical culture medium microvolume detection device 64 for detecting the presence of a stem cell bearing culture medium microvolume within a predetermined segment of the transfer tube 54. The computer 53 also controls a lifting device 66 for controlling the height of a source of stem cells 67 for selectively immersing the uptake catheter's distal end 59B therein, and a micromanipulator 68 for controlling the location of the delivery catheter's distal end 61B.

Operation of apparatus 51 for transplanting a single stem cell bearing flattened droplet at the target site is now described with reference to Figures 6A-6C is as follows:

The computer 53 sets the 2-way valve 58 to connect the uptake catheter 59 to the microvolume pump 56, operates the lifting device 66 to lift the stem cell source 67 to immerse the uptake catheter's distal end 59B therein, controls the microvolume pump 56 for drawing an incoming flow of displacement gas thereinto so as to aspirate a stem cell bearing culture medium microvolume into the uptake catheter 59, and then operates the lifting device 66 a second time to lower the stem cell source 67 to remove the distal end 59B therefrom (see Figure 6A).

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The computer 53 continues to operate the microvolume pump 56 to draw an incoming flow of displacement gas thereinto to aspirate the stem cell bearing culture medium microvolume into the transfer tube 57 towards its proximal end 57A. Upon its reaching the culture medium microvolume detection device 64, the
5 culture medium microvolume detection device 64 sends a signal to the computer 53 which interrupts the operation of the microvolume pump 56 and opens the NC venting valve 63 to stop the inward movement of the stem cell bearing culture medium microvolume (see Figure 6B).

The computer 53 closes the NC venting valve 63, sets the 2-way valve 58 to
10 connect the transfer tube 57 to the delivery catheter 61, and operates the microvolume pump 56 to issue an outgoing flow of displacement gas to outwardly displace the stem cell bearing culture medium microvolume towards the distal end 61B (see Figure 6C). The computer 53 operates the microvolume pump 56 to deposit a stem cell bearing flattened droplet at the target site in a similar manner as
15 described hereinabove with respect to the flattened droplet type IVF-ET procedure. The computer 53 operates the micromanipulator 68 to withdraw the delivery catheter's distal end 61B by about 0.5-3 mm to detach a stem cell bearing flattened droplet therefrom. The computer 53 also operates the micromanipulator 68 to gently manipulate the delivery catheter 61 on detection of a fault signature FS2 and
20 the microvolume pump 56 on detection of a fault signature FS3.

Figure 7 shows a Multiple Flattened Droplet Pressure Waveform (MFDPW) corresponding to the transportation of successive stem cell bearing flattened droplets to a target site. The MFDPW has a stepped appearance reflecting the fact that each transplantation of a stem cell flattened droplet at the target site increases
25 the volume thereat which in turn increases the prevailing pressure in the transfer tube 57 required to deposit the next flattened droplet. The MFDPW is effectively constituted by a series of SFDPWs (see Figure 4) except with an additional sub-atmospheric pressure segment between each pair of consecutive SFDPWs. The sub-atmospheric pressure segments each correspond to the loading of the uptake

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catheter 59 with a stem cell culture medium microvolume and thereafter its inward displacement to the transfer tube 57.

Figure 8 shows that stem cell bearing flattened droplets D tend to fuse into a single large drop L if deposited sufficiently adjacent to one another at the target
5 site.

Various modifications and changes may be made in the configuration described hereinabove that come within the spirit of the invention. The invention embraces all such changes and modifications coming within the scope of the
10 appended claims.